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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
09/373,230	08/12/1999	HARUKI OKMURA	OKAMURA=2E	2359	
	7590 08/08/2007 D NEIMARK, P.L.L.C.		EXAMINER		
624 NINTH ST	•		JIANG, DONG		
SUITE 300 WASHINGTO	N, DC 20001-5303		ART UNIT	PAPER NUMBER	
			1646		
			MAIL DATE	DELIVERY MODE	
•			08/08/2007	PAPER	

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Office Action Summary		Applicat	Application No. Applicant(s)					
		09/373,2	230	OKMURA ET AL.				
		Examine	er	Art Unit				
	·	Dong Jia	_ <del>_</del>	1646				
- Period fo	- The MAILING DATE of this communicati Reply	ion appears on ti	he cover sheet w	ith the correspondence ac	ldress			
WHICI - Extens after S - If NO   - Failure Any re	PRTENED STATUTORY PERIOD FOR HEVER IS LONGER, FROM THE MAIL sions of time may be available under the provisions of 37 kl X (6) MONTHS from the mailing date of this communical period for reply is specified above, the maximum statutory to reply within the set or extended period for reply will, by ply received by the Office later than three months after the patent term adjustment. See 37 CFR 1.704(b).	ING DATE OF T CFR 1.136(a). In no ention. by period will apply and soy statute, cause the apply	HIS COMMUNI event, however, may a will expire SIX (6) MON oplication to become A	CATION. reply be timely filed  NTHS from the mailing date of this c BANDONED (35 U.S.C. § 133).	,			
Status								
1)[🛛	Responsive to communication(s) filed or	n 05 June 2007						
		This action is		·				
·	Since this application is in condition for allowance except for formal matters, prosecution as to the merits is							
	closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11, 453 O.G. 213.							
	on of Claims	•	,	•	•			
4) 🖂 (	Claim(s) <u>18-23</u> is/are pending in the app	lication.	•	•				
	4a) Of the above claim(s) is/are withdrawn from consideration.							
	5) Claim(s) is/are allowed.							
6)🛛 (	)⊠ Claim(s) <u>18-23</u> is/are rejected.							
7) 🗌 (	Claim(s) is/are objected to.							
8) 🗌 (	Claim(s) are subject to restriction	and/or election	requirement.					
Application	on Papers							
9) <u></u> ⊤	he specification is objected to by the Ex	aminer.						
	he drawing(s) filed on is/are: a)[		o) ☐ objected to	by the Examiner.				
	Applicant may not request that any objection			•				
	Replacement drawing sheet(s) including the				FR 1.121(d).			
_	he oath or declaration is objected to by				• •			
Priority u	nder 35 U.S.C. § 119							
_	cknowledgment is made of a claim for for following the complex of	oreign priority ur	nder 35 U.S.C. §	§ 119(a)-(d) or (f).				
•	Certified copies of the priority docu	uments have be	en received.					
2	2. Certified copies of the priority docu	uments have be	en received in A	application No				
3	3. Copies of the certified copies of th	e priority docum	ents have been	received in this National	Stage			
	application from the International E	Bureau (PCT Ru	ıle 17.2(a)).		_			
* Se	ee the attached detailed Office action for	a list of the cer	tified copies not	received.				
Attachment(								
·	of References Cited (PTO-892) of Draftsperson's Patent Drawing Review (PTO-9	48)	=	Summary (PTO-413) s)/Mail Date				
3) 🔲 Informa	ation Disclosure Statement(s) (PTO/SB/08)		5) Notice of I	nformal Patent Application				
Paper	No(s)/Mail Date		6)	·····•				

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#### **DETAILED OFFICE ACTION**

The request filed on 05 June 2007 for a Continued Examination (RCE) under 37 CFR 1.114 based on parent Application No. 09/373,230 is acceptable, and a RCE has been established. An action on the RCE follows.

Applicant's amendment filed on 04 January 2007 is acknowledged and entered. Following the amendment, claims 3-9, 11, 14 and 16 are canceled, and the new claims 18 and 19 are added.

Applicant's amendment filed on 05 June 2007 is acknowledged and entered. Following the amendment, claims 18 and 19 are amended, and the new claims 20-23 are added.

Currently, claims 18-23 are pending and under consideration.

#### Withdrawal of Objections and Rejections:

All objections and rejections of claims 3-9, 11, 14 and 16 are moot as the applicant has canceled the claims.

#### New Matter Rejection

The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

Claims 20 and 21 are rejected under 35 U.S.C. 112, first paragraph, as containing subject matter which was not described in the specification in such a way as to reasonably convey to one skilled in the relevant art that the inventor(s), at the time the application was filed, had possession of the claimed invention. Applicants have not pointed out, nor can the Examiner locate, the basis in the specification for the limitation "one or more amino acids but *not so many*" in claim 20 (lines 6, 7 and 9).

This is a new matter rejection.

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#### Rejections under 35 U.S.C. 112:

The following is a quotation of the second paragraph of 35 U.S.C. 112:

The specification shall conclude with one or more claims particularly pointing out and distinctly claiming the subject matter which the applicant regards as his invention.

Claims 20-23 are rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

Claim 20 is indefinite for the recitation "one or more amino acids but *not so many*" in lines 6, 7 and 9 because it is unclear what it is meant by "not so many", and how many is not so many. The metes and bounds of the claim, therefore, cannot be determined.

Claim 22 is indefinite for the recitation "not identical to a protein" in line 2 because it is unclear what it is meant by "not identical", and how different is not identical. The metes and bounds of the claim, therefore, cannot be determined.

The remaining claims are included in this rejection because they are dependent from the specifically mentioned claims without resolving the indefiniteness issue belonging thereto.

The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

Claims 18-23 are rejected under 35 U.S.C. 112, first paragraph, as containing subject matter which was not described in the specification in such a way as to reasonably convey to one skilled in the relevant art that the inventor(s), at the time the application was filed, had possession of the claimed invention.

Claims 18-23 are directed to an variant of IL-18, which encompasses variants having one or more amino acids in SEQ ID NO:2 replaced (claim 18, part (i), for example), one or more amino acids but not so many replaced (claim 20, part (i), for example), and variants not identical to SEQ ID NO:2 (claim 22, for example), which read on functional equivalents of SEQ ID NO:2 as there is no clear limitation as to how many amino acids can be replaced in SEQ ID NO:2.

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Thus, the claims encompass significant structural dissimilarity as compared to the exemplified IL-18 and the variants, including functional equivalents without any sequence similarity to the disclosed SEQ ID NO:2 (claim 18, part (i), for example). However, the specification discloses one IL-18 amino acid sequences with particularity, the murine IL-18 with SEQ ID NO:2, and no particular variant of the IL-18 with amino acid addition, deletion, substitution, or any other type of "functional equivalents" meeting the limitations of these claims were ever identified or particularly described.

To provide adequate written description and evidence of possession of a claimed genus, the specification must provide sufficient distinguishing identifying characteristics of the genus. The factors to be considered include disclosure of compete or partial structure, physical and/or chemical properties, functional characteristics, structure/function correlation, methods of making the claimed product, or any combination thereof. In this case, the only factors present in the claims are MW and a functional characteristic, inducing IFN-γ production. There is no sequence similarity required for the claimed variants. Thus, With the exception of SEQ ID NO:2, the skilled artisan cannot envision the detailed chemical structure of the encompassed variants, and therefore conception is not achieved regardless of the complexity or simplicity of the method of making a peptide or chemical molecule. Accordingly, the specification does not provide adequate written description of the claimed genus.

<u>Vas-Cath Inc. v. Mahurkar</u>, 19USPQ2d 1111, clearly states that "applicant must convey with reasonable clarity to those skilled in the art that, as of the filing date sought, he or she was in possession of the invention. The invention is, for purposes of the 'written description' inquiry, whatever is now claimed." (See page 1117.) The specification does not "clearly allow persons of ordinary skill in the art to recognize that [he or she] invented what is claimed." (See *Vas-Cath* at page 1116).

Adequate written description requires more than a mere statement that it is part of the invention and reference to a potential method of isolating it. The compound itself is required. See Fiers v. Revel, 25 USPQ2d 1601 at 1606 (CAFC 1993) and Amgen Inc. v. Chugai Pharmaceutical Co. Ltd., 18 USPQ2d 1016.

One cannot describe what one has not conceived. See *Fiddes v. Baird*, 30 USPQ2d 1481 at 1483. In *Fiddes*, claims directed to mammalian FGF's were found to be unpatentable due to

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lack of written description for that broad class. The specification provided only the bovine sequence.

Therefore, only the polypeptide of SEQ ID NO:2, but not the full breadth of the claims meet the written description provision of 35 U.S.C. §112, first paragraph. Applicant is reminded that *Vas-Cath* makes clear that the written description provision of 35 U.S.C. §112 is severable from its enablement provision (see page 1115).

## **Conclusion:**

No claim is allowed.

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### **Advisory Information:**

Any inquiry concerning this communication should be directed to Dong Jiang whose telephone number is 571-272-0872. The examiner can normally be reached on Monday - Friday from 9:30 AM to 7:00 PM.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Gary Nickol, can be reached on 571-272-0835. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Dong Jiang, Ph.D. Patent Examiner AU1646

8/2/07